TYROBP gene

TYRO protein tyrosine kinase binding protein

Normal Function

The *TYROBP* gene (also known as the *DAP12* gene) provides instructions for making a protein called the TYRO protein tyrosine kinase binding protein. This protein is found in a variety of cells produced in bone marrow (myeloid cells) and other immune system cells (lymphoid cells). The protein is located on the cell surface, where it helps transmit chemical signals that activate the cell.

The TYROBP protein interacts with several other proteins on the surface of cells. For example, it forms a complex with the protein produced from the *TREM2* gene. The TYROBP protein and its partners were first identified in the immune system, where they activate certain cells (such as natural killer cells and dendritic cells) that trigger an inflammatory response to injury or disease.

The TYROBP-TREM2 complex also activates cells in the skeletal system and in the brain and spinal cord (central nervous system). In the skeletal system, the complex is found in osteoclasts, which are specialized cells that break down and remove (resorb) bone tissue that is no longer needed. These cells are involved in bone remodeling, which is a normal process that replaces old bone tissue with new bone. In the central nervous system, the complex appears to play an important role in immune cells called microglia. These cells protect the brain and spinal cord from foreign invaders and remove dead nerve cells and other debris. Although the TYROBP-TREM2 complex plays a critical role in osteoclasts and microglia, its exact function in these cells is unclear.

Health Conditions Related to Genetic Changes

polycystic lipomembranous osteodysplasia with sclerosing leukoencephalopathy

At least six mutations in the *TYROBP* gene have been identified in people with polycystic lipomembranous osteodysplasia with sclerosing leukoencephalopathy (commonly known as PLOSL). One *TYROBP* mutation has been found to cause PLOSL in all affected people of Finnish ancestry. This mutation deletes a significant portion of the *TYROBP* gene, which prevents the cell from producing any protein from this gene. Mutations in other populations result in the production of an abnormally short, nonfunctional version of the protein or prevent the protein from reaching the cell surface.

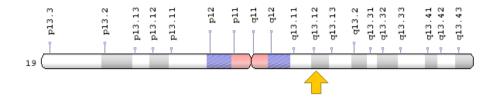
Researchers believe that the signs and symptoms of PLOSL are related to defective TYROBP-TREM2 signaling in osteoclasts and microglia. The bone abnormalities

seen with this disorder are probably related to malfunctioning osteoclasts, which are less able to resorb bone tissue during bone remodeling. In the central nervous system, defective signaling through the TYROBP-TREM2 complex causes widespread abnormalities of microglia. Researchers are working to determine how these abnormalities lead to the neurological problems associated with PLOSL.

Chromosomal Location

Cytogenetic Location: 19q13.12, which is the long (q) arm of chromosome 19 at position 13.12

Molecular Location: base pairs 35,904,401 to 35,908,309 on chromosome 19 (Homo sapiens Annotation Release 108, GRCh38.p7) (NCBI)



Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- DAP12
- DNAX-activation protein 12
- KAR-associated protein
- KARAP
- killer activating receptor associated protein
- TYOBP_HUMAN

Additional Information & Resources

Educational Resources

- Basic Neurochemistry (sixth edition, 1998): The microglial cell plays a role in phagocytosis and inflammatory responses https://www.ncbi.nlm.nih.gov/books/NBK28217/#A37
- Molecular Biology of the Cell (fourth edition, 2002): Bone is continually remodeled by the cells within it https://www.ncbi.nlm.nih.gov/books/NBK26889/#A4187

GeneReviews

 Polycystic Lipomembranous Osteodysplasia with Sclerosing Leukoencephalopathy (PLOSL)

https://www.ncbi.nlm.nih.gov/books/NBK1197

Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28TYROBP%5BTIAB%5D%29+OR+%28TYRO+protein+tyrosine+kinase+binding+protein%5BTIAB%5D%29%29+OR+%28DAP12%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+2880+days%22%5Bdp%5D

OMIM

 TYRO PROTEIN TYROSINE KINASE-BINDING PROTEIN http://omim.org/entry/604142

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology http://atlasgeneticsoncology.org/Genes/GC_TYROBP.html
- ClinVar https://www.ncbi.nlm.nih.gov/clinvar?term=TYROBP%5Bgene%5D
- HGNC Gene Symbol Report http://www.genenames.org/cgi-bin/gene_symbol_report?q=data/ hqnc data.php&hqnc id=12449
- NCBI Gene https://www.ncbi.nlm.nih.gov/gene/7305
- UniProt http://www.uniprot.org/uniprot/O43914

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 GeneReview: Polycystic Lipomembranous Osteodysplasia with Sclerosing Leukoencephalopathy (PLOSL)

https://www.ncbi.nlm.nih.gov/books/NBK1197

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